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# Somatostatin Binding Capacity, Guanylate Cyclase and Tyrosine Phosphatase Activities During Pancreatic Proliferation in the Rat Induced by Gastrectomy

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RODRÍGUEZ-MARTÍN, E., A. M. VALENCIA, B. COLÁS, C. GARCÍA-ESCRIBANO, M. RODRÍGUEZ-PUYOL, C. SUSINI AND E. ARILLA. *Somatostatin binding capacity, guanylate cyclase and tyrosine phosphatase activities during pancreatic proliferation in the rat induced by gastrectomy*. PEPTIDES 16(8) 1461–1467, 1995. —Gastrectomy increased pancreatic growth and this effect was associated with an increase in the number of somatostatin-14 (SS) receptors (146% of control) without altering their affinity. SS increased guanylate cyclase activity twofold in pancreatic acinar membranes from gastrectomized rats. The gastrectomy decreased pancreatic SS-like immunoreactivity (SS-LI) content (55% of control levels) and tyrosine phosphatase activity (74% of control levels). Administration of proglumide (20 mg/kg, IP), a gastrin/cholecystokinin (CCK) receptor antagonist, suppressed the inhibitory effect of gastrectomy on basal tyrosine phosphatase activity and SS-LI content, which returned to control levels. Furthermore, proglumide suppressed the increase of the number of SS receptors and of SS-stimulated guanylate cyclase activity induced by gastrectomy. All this suggests that pancreatic acinar cell growth is associated with upregulation of SS receptors, which could represent a mechanism promoted by the cell to negatively regulate the mitogenic activity of pancreatic growth factors such as CCK. In addition, the results also suggest that the negative regulation of tyrosine phosphatase activity may be important in the events involved in the pancreatic hyperplasia observed after gastrectomy.

Gastrectomy	Guanylate cyclase	Somatostatin receptors	Pancreatic acinar membranes
Somatostatin like-immunoreactivity		Rat	

SOMATOSTATIN-14 (SS) is a tetradecapeptide synthesized in endocrine D cells that is widely distributed in the gastrointestinal tract (4,21) and is known to inhibit the stimulatory effects of various secretagogues on a number of cell functions, including endocrine and exocrine functions of the pancreas (53). More recently, SS analogues have been shown to inhibit the growth of normal and tumoral cells (23,45). In the pancreas, SS inhibited normal (23,45) and caerulein-stimulated pancreatic growth (31,42), whereas the neutralization of the endogenous SS with antiserum amplified the growth-stimulating effect of caerulein (31). These observations have led to the suggestion that this peptide has a physiologically important function in the regulation of normal cell growth and that it might be a negative regulator of pancreatic acinar cell proliferation in vivo. SS exerts its physiological actions by interacting with membrane-bound receptors

(48,58), which are coupled to adenylate cyclase (43), K<sup>+</sup> channels (61), voltage-dependent Ca<sup>2+</sup> channels (62), and tyrosine phosphatase (12). In addition, several authors have shown that SS increases the guanylate cyclase activity in several tissues (11,17,54). Recent studies argue that, in contrast to cyclic AMP (57), cyclic GMP has a role in the antiproliferative effect of SS (41).

One strong stimulant of exocrine pancreatic growth in the rat is a total gastrectomy, which results in acinar cell hyperplasia 2 weeks after the intervention (10). To determine whether SS is involved in exocrine pancreas growth responses in vivo, we measured the status of SS binding and pancreatic somatostatin-like immunoreactivity (SS-LI) content in control pancreas and during pancreatic proliferation 2 weeks after gastrectomy in rats. This study also examines the basal activity of the guanylate cyclase

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system as well as the stimulatory effect of SS on this enzyme activity in these membranes. Cholecystokinin (CCK) is the major humoral candidate in promoting postgastrectomy pancreatic hyperplasia (9,29). Proglumide, an antagonist of the receptors of the gastrin/CCK peptide family (37), was used to evaluate whether the effects of gastrectomy on the pancreatic somatostatinergic system involved the activation of gastrin/CCK receptors. Protein tyrosine phosphatases have been reported to play a crucial role in cellular processes involving protein phosphorylation such as growth and differentiation (60). The fact that tyrosine phosphatase activity is stimulated by growth factor inhibitors, like SS (50) and transforming growth factor  $\beta$  (TGF- $\beta$ ) (20), and by differentiating agents (51,53) that inhibit cell growth suggests an inverse relationship between tyrosine phosphatase activity and cell growth. Therefore, protein tyrosine phosphatase activity was also determined in the membrane preparations previously cited.

#### METHOD

##### Materials

Synthetic [Tyr<sup>11</sup>]SS was purchased from Universal Biologicals Ltd. (Cambridge, UK); collagenase (from *Clostridium histolyticum*) was obtained from Serva Fine Chemicals (Tebu, France). Bacitracin, phenylmethylsulfonylfluoride (PMSF), guanosine triphosphate (GTP), 3-isobutyl-1-methylxanthine (IBMX), bovine serum albumin (BSA), Nonidet P-40, poly(Glu,Tyr) (4:1 by mass), ammonium molybdate (1% in 1.32 M HCl) were purchased from Sigma (St. Louis, MO). [ $\gamma$ -<sup>32</sup>P]ATP was from NEN (Les Ulis, France). Sephacryl S-300 HR was from Pharmacia (Uppsala, Sweden). Carrier-free Na[<sup>125</sup>I] (IMS 300, 100 mCi/ml), the rabbit antibody used in the radioimmunoassay technique, and the cGMP kit were purchased from the Radiochemical Center (Amersham, UK). This antiserum was raised in rabbits against SS conjugated to BSA and is specific for SS, but because SS constitutes the C-terminal portion of both somatostatin-25 (SS-25) and somatostatin-28 (SS-28), the antiserum does not distinguish between these three forms. The binding of SS to this antibody does not depend on an intact disulfide bond in the molecule, because breaking the disulfide bond by reaction with 0.1% mercaptoethanol (5 min boiling water bath) does not change peptide immunoreactivity. Cross-reactivity with other peptides was less than 0.5%. Cross-reaction with several SS analogues demonstrated that neither the N-terminal glycine nor the C-terminal cysteine residues are required for antibody binding, suggesting that the antigen site is directed towards the central part of the molecule containing the tryptophan residue. All other reagents were of the highest purity commercially available.

##### Experimental Animals

Male Wistar rats with an initial body weight of 200–250 g were used for the study. Animals were fasted for 12 h before surgery. Using ether anesthesia, total gastrectomy was performed through a midline incision by removing the entire stomach. The vagal trunks were preserved and continuity restored by end-to-end anastomosis between the esophagus and duodenum. Sham operation was performed by making and repairing an incision in the stomach. In the proglumide-treated groups, the animals received two times daily for 2 weeks an IP injection of proglumide (20 mg/kg). Proglumide was dissolved in saline solution. Postoperatively the rats were maintained on a 5% glucose diet for the first 24 h, then allowed free access to standard chow and tap water. These rats were sacrificed by decapitation 2 weeks after the intervention and the pancreas was removed and trimmed free of fat, connective tissues, and lymph nodes.

##### Preparation of Rat Pancreatic Acinar Membranes

Dispersed pancreatic acini were obtained from male Wistar rats after enzymatic degradation of the pancreas with 0.2 units of collagenase/ml in an oxygenated Krebs–Ringer medium as described by Amsterdam et al. (2). After washing by sedimentation, acini were transferred to 0.3 M sucrose, 0.03% soybean trypsin inhibitor, 0.1 mM PMSF, and 1 mM benzamidine (buffer A). In this buffer, the acini were homogenized at 4°C by use of a Potter homogenizer. After centrifugation at 600  $\times$  g for 5 min at 4°C, the supernatant was resuspended in buffer A and centrifuged at 100,000  $\times$  g in a TLA 100.3 rotor for 1 h. The supernatant was discarded and the pellet was resuspended in 50 mM Tris-HCl, pH 7, 0.5 mg/ml bacitracin, 0.2 mM CaCl<sub>2</sub>, 0.03% soybean trypsin inhibitor, 0.1 mM PMSF, 1 mM benzamidine, and 5% glycerol (buffer B). An aliquot was taken for protein determination by the method of Lowry et al. (27) with bovine serum albumin as a standard.

##### Binding of [<sup>125</sup>I-Tyr<sup>11</sup>]SS

Binding of [<sup>125</sup>I-Tyr<sup>11</sup>]SS was assayed on pancreatic acinar membranes from Wistar rats by a modified method (12). [<sup>125</sup>I-Tyr<sup>11</sup>]SS was radioiodinated by chloramine-T iodination according to the method of Greenwood (19). Separation of iodinated SS from unincorporated iodine was carried out on a Sephadex G-25 (fine) column equilibrated and eluted with 0.1 M acetic acid in BSA (0.1% w/v). The specific activity of the radioligand was 600 Ci/mmol.

Binding of [<sup>125</sup>I-Tyr<sup>11</sup>]SS to pancreatic acinar membranes was carried out in a total volume of 250  $\mu$ l in 50 mM Tris-HCl buffer (pH 7.4) containing 0.5 mM MgCl<sub>2</sub>, 3 mM NaCl, 0.2 mM CaCl<sub>2</sub>, 0.2% (w/v) BSA, 0.5 mg/ml bacitracin, and 0.3 mg/ml soybean trypsin inhibitor (binding buffer). Plasma membranes (140  $\mu$ g of protein/ml) were incubated for 90 min at 20°C with 100 pM [<sup>125</sup>I-Tyr<sup>11</sup>]SS in the absence or presence of unlabeled SS. Bound and free ligand was separated by centrifugation at 11,000  $\times$  g for 4 min at 4°C in a microcentrifuge. Radioactivity in the pellet was measured with a gamma scintillation counter. Nonspecific binding was estimated as membrane-associated radioactivity in the presence of 1  $\mu$ M SS and specific binding was calculated as the difference between total and nonspecific membrane-associated radioactivity.

##### Evaluation of Radiolabeled Peptide Degradation

To determine the extent of tracer degradation during incubation, we measured the ability of preincubated peptide to bind to fresh pancreatic acinar membranes. Briefly, [<sup>125</sup>I-Tyr<sup>11</sup>]SS (100 pM) was incubated with pancreatic acinar membranes (140  $\mu$ g protein/ml) for 90 min at 20°C. After this preincubation, aliquots of the medium were added to fresh pancreatic acinar membranes and incubated for an additional 90 min at 20°C. The fraction of the added radiolabeled peptide that specifically bound during the second incubation was measured and expressed as a percentage of the binding that was obtained in control experiments performed in the absence of pancreatic acinar membranes during the preincubation period. Pancreatic acinar cell membranes from all the groups showed a similar peptide degradation capacity and the values varied by no more than 9% in all the experimental groups.

##### Measurement of the Particulate Guanylate Cyclase Activity

The activity of particulate guanylate cyclase in pancreatic acinar membranes was measured using the method previously described (16,25) with a slight modification. The membranes (50  $\mu$ g protein/10  $\mu$ l) were resuspended in 50 mM Tris-HCl, pH 7.6,

and used for the assay of guanylate cyclase activity. A 10- $\mu$ l aliquot of the particulate fraction was added to tubes containing 80  $\mu$ l of a reaction mixture containing (final concentrations) 50 mM Tris-HCl, pH 7.6, 40 mM MnCl<sub>2</sub>, 1 mM GTP, 1 mM IBMX, 0.01% bacitracin, and a GTP-regenerating system that consisted of 15 mM creatine phosphate and 20  $\mu$ g of creatine phosphokinase. After incubation at 37°C for 10 min with or without SS ( $10^{-7}$ – $10^{-12}$  M) the reaction was stopped by the addition of 900  $\mu$ l of 50 mM sodium acetate, pH 4.0, followed by heating at 90°C for 3 min. The heated incubation mixture was centrifuged at  $2000 \times g$  for 20 min and the supernatant fraction was evaporated to dryness. The amount of cyclic GMP generated was determined in the dry extract by radioimmunoassay.

#### Tissue Extraction and SS Radioimmunoassay

For SS-LI measurement, the pancreata were rapidly homogenized in 1 ml 2 M acetic acid using a Brinkman polytron (setting 5, 30 s). The extracts were boiled for 5 min in an ice-chilled water bath, and aliquots (100  $\mu$ l) were removed for protein determination (27). The homogenates were subsequently centrifuged at  $15,000 \times g$  for 15 min at 4°C and the supernatant was neutralized with 2 M NaOH. The extracts were stored at –70°C until assay. The SS concentration was determined in tissue extracts by a modified radioimmunoassay method (36), with a sensitivity limit of 10 pg/ml. Incubation tubes prepared in duplicate contained 100- $\mu$ l samples of tissue extracts or standard solutions of 0–500 pg cyclic SS diluted in phosphate buffer (0.05 M, pH 7.2, containing 0.3% BSA, 0.01 M EDTA), 200  $\mu$ l appropriately diluted anti-SS-serum, 100  $\mu$ l freshly prepared [<sup>125</sup>I-Tyr<sup>11</sup>]SS diluted in buffer to give 6000 cpm (equivalent to 5–10 pg), in a final volume of 0.8 ml. All reagents as well as the assay tubes were kept chilled on ice before incubation for 48 h at 4°C. Bound hormone was separated from free hormone by the addition of 1 ml dextran-coated charcoal (dextran T-70, 0.2% w/v, Pharmacia; charcoal: Norit A, 2% w/v, Serva, Feinbiochemica, Heidelberg, Germany). Serial dilution curves for the samples were parallel with the standard curve. The intra- and interassay variation coefficients were 6.0% and 8.8%, respectively.

#### Preparation of A431 Membranes

A431 cells were washed once with 10 mM sodium phosphate, pH 7, and 150 mM NaCl (NaCl/P) and scraped into 10 mM NaCl/P, pH 7 (18). The cells were collected by centrifugation, suspended in 50 mM Tris-HCl, pH 7, containing 0.03% soybean trypsin inhibitor, 0.5 mg/ml bacitracin, and lysed by freeze-thawing in liquid nitrogen. The lysate was centrifuged at  $170 \times g$  for 5 min. The subsequent supernatant was centrifuged at  $25,000 \times g$  for 30 min. The final pellet was resuspended in the 50 mM Tris-HCl, pH 7, containing 0.03% soybean trypsin inhibitor and 0.5 mg/ml bacitracin (12).

#### Preparation of Phosphorylated Poly(Glu,Tyr)

Phosphorylated poly(Glu,Tyr) was prepared by incubating 7.2 mg polymer with 1.5–2 mg A431 cell membranes in 50 mM Tris-HCl, containing 0.1  $\mu$ M epidermal growth factor, 250  $\mu$ M [ $\gamma$ -<sup>32</sup>P]ATP (700 dpm/pmol), 100  $\mu$ M orthovanadate, 5 mM MnCl<sub>2</sub>, and 0.1% Nonidet P-40 at pH 7.5 (12). The reaction was allowed to proceed for 16 h at 4°C, then the mixture was subjected to gel filtration on a Sephacryl S300 HR column. Fractions containing the phosphorylated poly(Glu,Tyr) were pooled and concentrated in a Centricon-3 Amicon microconcentrator; 100- $\mu$ l aliquots were stored at –20°C.

#### Assay for Phosphotyrosine Phosphatase Activity

PTPase activity was measured by the release of [<sup>32</sup>P]orthophosphate from <sup>32</sup>P-labeled poly(Glu,Tyr) in a 100- $\mu$ l reaction mixture containing 30,000 cpm <sup>32</sup>P-labeled poly(Glu,Tyr) (final concentration of 0.3  $\mu$ M) and 3  $\mu$ g acinar pancreatic membrane proteins in buffer B containing 0.1% albumin. The reaction was allowed to proceed for 5 min at 30°C, then stopped by the addition of 100  $\mu$ l 30% trichloroacetic acid. The mixtures were kept on ice for 30 min, then centrifuged at  $10,000 \times g$  for 10 min to remove the denatured proteins. The inorganic [<sup>32</sup>P]phosphate liberated was extracted using the molybdate extraction procedure (3) and radioactivity was evaluated by liquid scintillation. The amount of <sup>32</sup>P<sub>i</sub> released was determined from the specific radioactivity of [ $\gamma$ -<sup>32</sup>P]ATP used for the phosphorylation reaction. One unit PTPase activity was defined as the amount that released 1 nmol phosphate/min at 30°C from radiolabeled substrate.

#### Data Analysis

The computer program LIGAND (34) was used to analyze the binding data. The use of this program made it possible to select the receptor models that best fit a given set of binding data. The same program was also used to present data in the form of Scatchard plots (44) and to compute the values for receptor affinity ( $K_d$ ) and density ( $B_{max}$ ) that best fit the sets of binding data for each rat. Statistical comparisons of all the data were analyzed by ANOVA and the Newman-Keuls *t*-test. Means among groups were considered significantly different when the value was  $p < 0.05$ . Each individual experiment was performed in duplicate.

#### RESULTS

Two weeks following total gastrectomy, the weight of the pancreas increased significantly by 30% ( $p < 0.001$ ) when compared to sham-operated rats. Pancreatic acinar membrane PTPase activity decreased (74% of control) 2 weeks after total gastrectomy, coinciding with hyperplasia of exocrine pancreatic tissue previously described (Fig. 1).

Figure 2 shows that pancreatic SS-LI content 2 weeks after total gastrectomy was significantly lower (55% of control levels) than in control rats.

The specific binding of SS to rat pancreatic acinar membranes increased (146% of control) after gastrectomy compared to con-

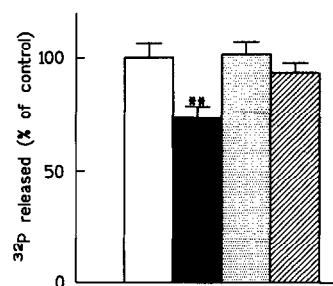


FIG. 1. Rat pancreatic acinar membrane phosphotyrosine phosphatase (PTPase) activity. Crude membranes were prepared as described in the Method section. PTPase activity was expressed as a percentage of control activity obtained with pancreatic acinar membranes from sham-operated rats. Control values averaged  $0.13 \pm 0.009$  nmol/mg protein/min. Results are means  $\pm$  SEM of five different experiments performed in duplicate. \*\* $p < 0.01$  vs. respective control. Open bar: sham-operated; black bar: gastrectomized; speckled bar: gastrectomized plus proglumide; hatched bar: proglumide alone.

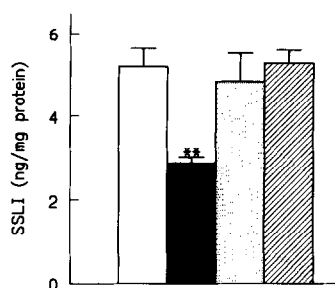


FIG. 2. Pancreatic somatostatin-like immunoreactivity (SS-LI) content in sham-operated (open bar), gastrectomized (black bar), gastrectomized plus proglumide-treated (speckled bar), and proglumide-treated (hatched bar) rats. Values are expressed as the mean  $\pm$  SEM in ng SS/mg protein for five rats per group performed in duplicate. Statistical comparison vs. control: \*\* $p < 0.01$ .

tol conditions. When comparing the corresponding curves of [ $^{125}$ I-Tyr $^{11}$ ]SS displacement by increasing concentrations of the unlabeled peptide (Fig. 3), the binding data were significantly higher in the gastrectomized group throughout the whole range studied. Scatchard plots of the stoichiometric binding data were linear and essentially parallel (Fig. 3, right panel). Interpretation of these data with the LIGAND computer program (34) resulted in the best fit for a model with one type of SS receptor. Pancreatic acinar membranes from gastrectomized rats exhibited significant increases in the maximum SS binding capacity. However, the corresponding  $K_d$  values remained unchanged after gastrectomy.

The modulation of guanylate cyclase by SS was studied in pancreatic acinar membranes from control and gastrectomized rats. No significant differences were seen for basal guanylate cyclase activities between the control group and the gastrectomized group. The capacity of SS to stimulate guanylate cyclase activity in control and gastrectomized rats is shown in Table 2. SS increased by twofold guanylate cyclase activity in pancreatic membranes from gastrectomized rats.

Because the major humoral candidate for promoting pancreatic hyperplasia after gastrectomy appears to be CCK (9,29), a group of gastrectomized rats was treated with proglumide, an antagonist of receptors of the gastrin/CCK peptide family to evaluate whether the effects of gastrectomy on the pancreatic somatostatinergic system involved the activation of gastrin/CCK. Proglumide suppressed the inhibitory effect of gastrectomy on basal tyrosine phosphatase activity and SS-LI content which returned to control levels (Figs. 1 and 2). Furthermore, proglumide suppressed the increase of the number of SS receptors and that of guanylate cyclase activity induced by gastrectomy (Fig. 3, Tables 1 and 2). Proglumide was also administered to sham-operated control rats without producing significant differences with the controls in any of the studied parameters.

#### DISCUSSION

Gastrectomy increased pancreatic growth (10) and this effect was associated with an increase in the number of SS receptors without alteration of affinity.

The molecular basis for the increased number of SS receptors after gastrectomy is uncertain. Because growth factor-induced cell division is paralleled by heteroregulation of other cell surface receptors (7), it is possible that SS receptors are targets for heterologous regulation by growth factors. Thus, the increase of SS receptors could be secondary to the CCK mitogenic signal (13,32). In this regard, the CCK receptors seem to mediate the action of gastrectomy on the SS receptor-effector system in the

pancreatic acinar membranes because the gastrectomy-induced changes in the somatostatinergic system were prevented by the gastrin/CCK receptor antagonist proglumide. In addition, proglumide alone had no demonstrable effect on these parameters. Two types of receptors for the gastrin/CCK peptide family have been identified in pancreatic acinar cells (37,59): the CCK $_A$  receptors, which have a much higher affinity for CCK than for gastrin (37); and the gastrin receptors (CCK $_B$  or GR), which have approximately the same affinity for CCK and gastrin (37).

The increase in the number of SS receptors observed during pancreatic acinar cell hyperplasia 2 weeks after total gastrectomy is consistent with the view that SS may function as a negative regulator of pancreatic acinar cell proliferation *in vivo* (31,42,46). Pancreatic acinar cells, which do not synthesize SS but possess functional receptors for SS (48,58), may respond to SS produced in the pancreas by the D cells in the islets of Langerhans as well as to circulating SS.

Recently we found that exocrine pancreatic growth observed after proximal small bowel resection is accompanied by an increase in the number of SS receptors at 2 weeks and 1 month after intestinal surgery (1). In addition, in pancreatic tumor cells AR42J, we also observed an increase in the number of SS receptors after treatment of cells with gastrin and epidermal growth factor (EGF) (and with CCK, not published), which stimulate cell growth (55). In contrast, conditions that decrease cell growth, such as treatment with glucocorticoids, decrease the number of SS receptors on AR42J cells (56). All these results suggest that pancreatic acinar cell growth is associated with upregulation of SS receptors and this could represent a mechanism promoted by the cell to negatively regulate the mitogenic activity of pancreatic growth factors such as CCK.

Although at least five SS receptor subtypes have been cloned (6), only SSTR3 (63) and SSTR5 (38) are present in the islets of Langerhans whereas the exocrine pancreas appears to express only receptor subtype 2 (SSTR2) (8). Vidal et al. (55) also reported that AR42J cells expressed SSTR2 subtype strongly and that the gastrin- and EGF-induced increase in SS receptors was due to upregulation in SSTR2 subtype receptors. All these results suggest that the increase in SS receptor number after gastrectomy

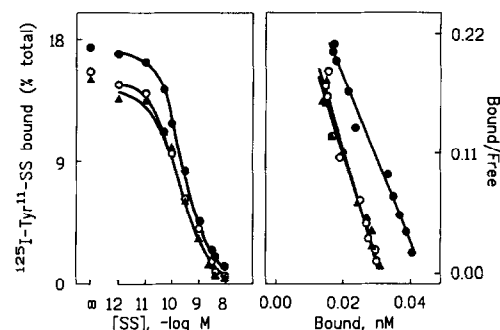


FIG. 3. Left panel: competitive inhibition of specific [ $^{125}$ I-Tyr $^{11}$ ]-somatostatin ([ $^{125}$ I-Tyr $^{11}$ ]SS) binding to rat pancreatic acinar membranes by unlabeled somatostatin. Points correspond to sham-operated ( $\circ$ ), gastrectomized ( $\bullet$ ), and gastrectomized plus proglumide ( $\blacktriangle$ ) rats. Each point is the mean of five separate experiments, each performed in duplicate. Results express the value of a pool of control groups because maximal binding capacity and dissociation constant values of the control groups were not affected. For the sake of clarity, SEM are not represented but were always below 10% of mean values. Right panel: Scatchard analysis of the same data. The kinetic constants calculated by Scatchard analysis are given in Table 1.

TABLE 1

EQUILIBRIUM PARAMETERS FOR SOMATOSTATIN (SS) BINDING TO PANCREATIC ACINAR MEMBRANES OF RATS THAT UNDERWENT A SHAM OPERATION, TOTAL GASTRECTOMY, TOTAL GASTRECTOMY PLUS PROGLUMIDE ADMINISTRATION, OR PROGLUMIDE ADMINISTRATION ALONE

Groups	SS Receptors	
	$B_{max}$ (fmol/mg protein)	$K_d$ (mM)
Sham operated	216 ± 21	0.09 ± 0.007
Gastrectomized	316 ± 30*	0.13 ± 0.008
Gastrectomized + proglumide	222 ± 17	0.10 ± 0.006
Proglumide	218 ± 13	0.09 ± 0.008

Binding parameters were obtained by computer nonlinear regression analysis (LIGAND) of individual competitive binding curves. Each value is the mean ± SEM of five separate experiments performed in duplicate. Comparison of means within each parameter group was by one-way ANOVA and Newman-Keuls tests.

\* Statistical comparison versus control:  $p < 0.01$ .

could result, at least in part, from upregulation of the SSTR2 subtype.

With respect to postreceptor mechanisms, recently Viguerie et al. (57) and Alvaro-Alonso et al. (1) reported that the cyclic AMP pathway is not involved in the inhibitory effect of SS of pancreatic cell growth. Because recent papers support a role for cyclic GMP in the antiproliferative effect of SS (41), the present study has examined the effect of SS on guanylate cyclase activity in pancreatic acinar cell membranes from control and gastrectomized rats. The cyclic GMP levels in pancreatic acinar membranes obtained in the present study are similar to those obtained by García-Escribano et al. (17) in mesangial cells and by Catalan et al. (11) in the cerebral cortex, and lower than those obtained by Vesely et al. (54) in the pancreas. However, the two latter authors measured the guanylate cyclase activity in the supernatant, whereas our study and that of García-Escribano et al. (17) measured the guanylate cyclase activity in the pellet. The present results show that SS increases guanylate cyclase activity significantly in the operated rats. However, the increase is not significant in the control rats, where it is comparable to the effect of proglumide alone. The increase does not appear to be due to a change in the number of molecules of guanylate cyclase itself because the activity levels were similar in membranes from both

control and gastrectomized animals. The intensity of the stimulatory effect of SS on guanylate cyclase activity in the gastrectomized animals is most probably related to the observed increase in SS receptors. These results demonstrate for the first time that SS stimulates membrane guanylate cyclase activity in pancreatic acinar cells. This effect is observed at a SS concentration that correlates with the affinity of SS for pancreatic SS receptors (22) and could be involved in the inhibitory effect of SS on pancreatic cell growth.

In addition to the intestinal mucosa, which preferentially releases SS-28 (5), a major site of endogenous SS release is located within the pancreas itself, namely in the D cells of the islets of Langerhans. The close interaction between the islets and the exocrine tissue appears to be at least partially mediated by an insuloacinar portal system (26,35). Thus, because of the relatively short plasmatic half-life of SS, which has been estimated to be approximately 1–2 min for SS and approximately 4 min for SS-28 (52), it seems that islet-released SS might act via the insuloacinar portal system (33). Hence, despite low peripheral plasma levels, relatively high local plasma concentrations would be present around the acinar tissue and this probably has functional relevance (33). Therefore, the present study measured pancreatic SS-LI content. The pancreatic SS-LI levels in the control rats were similar to those previously reported by others (36). Gastrectomy decreased pancreatic SS-LI content compared with controls. These results also agree with other experimental models of pancreatic growth studied by our group. Thus, two weeks after enterectomy, the situation is similar: increased SS receptors and decreased SS-LI (albeit not significant after enterectomy). Total gastrectomy leads to increased CCK (29) and glucagon (14) plasma levels compared to controls. Because it has been demonstrated that CCK and glucagon increases SS secretion (30), the decrease of pancreatic SS-LI levels in response to gastrectomy could be a consequence of increased plasma levels of both hormones.

Cell membrane-associated tyrosine phosphatase activities have been characterized in various tissues (15,24). Recently, Colás et al. (12) demonstrated that the level of tyrosine phosphatase activity in rat pancreatic acinar cells is as high as that observed in some rich sources of tyrosine phosphatase such as rat spleen (49) or rabbit kidney (47). The control values of the present study are in agreement with previously reported values. Gastrectomy decreased tyrosine phosphatase activity. The regulation of protein phosphorylation level in tyrosine residues is well known to be involved in cellular activities associated with growth and dif-

TABLE 2

EFFECT OF SOMATOSTATIN (SS) ON GUANYLATE CYCLASE ACTIVITY IN PANCREATIC ACINAR MEMBRANES OF RATS THAT UNDERWENT A SHAM OPERATION, TOTAL GASTRECTOMY, TOTAL GASTRECTOMY PLUS PROGLUMIDE ADMINISTRATION, OR PROGLUMIDE ADMINISTRATION ALONE

[SS]	Sham-Operated	Gastrectomized	Gastrectomized + Proglumide	Proglumide
0	9.3 ± 1.5	9.8 ± 0.9	9.5 ± 0.7	9.8 ± 1.4
1 pM	10.8 ± 0.9	11.8 ± 1.7	10.8 ± 1.9	10.3 ± 0.03
10 pM	11.5 ± 0.5	13.8 ± 1.3	11.5 ± 1.3	11.1 ± 0.8
100 pM	13.8 ± 1.2	16.3 ± 1.8	13.3 ± 0.6	13.6 ± 1.2
1 nM	14.4 ± 1.2	19.2 ± 0.4*	14.3 ± 1.5	14.2 ± 1.6
10 nM	13.7 ± 1.0	16.9 ± 0.7†	13.4 ± 0.2	13.5 ± 1.4
100 nM	12.6 ± 0.8	13.0 ± 0.8	12.4 ± 0.2	12.5 ± 1.9

Values are means ± SEM of determinations performed in pmol cyclic GMP/mg protein/10 min of incubation. Experiments were performed as described in the Method section.

\* † Statistical comparison vs. control: \* $p < 0.01$ , † $p < 0.05$ .

ferentiation. Indeed, it has been demonstrated that CCK stimulated tyrosine kinase activity in pancreatic acini (28) and this stimulation is associated with the growth-promoting effect of caerulein (40). Conversely, SS stimulation of membrane tyrosine phosphatase activity has been shown to be involved in the antiproliferative effect of SS in pancreatic tumor cells (50). The decrease of tyrosine phosphatase activity observed after gastrectomy is like that observed after pancreatectomy during the regeneration process associated with pancreatic growth (40). These observations suggest that the negative regulation of tyro-

sine phosphatase activity may be important in the events involved in the pancreatic hyperplasia observed after gastrectomy.

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